(35 minutes v 195 minutes respectively), a finding which might be explained by the differences in absorption of apomorphine from the two preparations.

These preliminary results suggest that apomorphine is effective when administered rectally as an enema and may be the chosen method for rapid relief of "off period" symptoms when other routes may be unavailable due to adverse effects or the patients' difficulty in applying subcutaneous or intranasal apomorphine. We advise regular proctoscopy when treatment is frequent or prolonged.

T VAN LAAR
ENH JANSEN
Department of Neurology,
The Netherlands
AWG ESSINK
WJ RUTTEN
C NEEF
Department of Clinical Pharmacy,
Hospital Medisch Spectrum Twente,
Emscherveld

Correspondence to: Dr Earnst Jansen, Department of Neurology and Movement Disorder Unit, Hospital Medisch Spectrum Twente, Haakberkersstraat 55, 7513 ER Enschede, The Netherlands.


HAMI TSP attributable to blood transfusion

A multi-centric case-control study was carried out to clarify possible environmental factors related to the onset of HTLV-I-associated myelopathy/tropical spastic paraparesis (HAM/TSP) in northern Kyushu, Japan, which comprises one of the most prevalent areas of HTLV-I in the world.

The frequencies of blood transfusion before the onset of HAMI TSP were 33% (6/18) among male patients and 18% (12/67) among female patients, which were significantly higher than 8% in males and 9% in females in the general population. The age-adjusted summary odds ratios with 95% confidence intervals were 7.0 (2.9-17.0) for males and 2.4 (1.3-4.5) for females. The percentages of population attributable risk of HAMI TSP attributable to transfusion were estimated to be approximately 29% (5-52) for males and 11% (1-21) for females.

The fraction of HAMI TSP attributable to transfusion after the introduction of blood screening for HTLV-I, in effect since 1986 in Japan, was definitely smaller than that before the programme. Our observations seemed compatible with a marked decline in newly diagnosed HAMI TSP patients after its introduction, which may be part of the benefit of blood screening. Neither smoking nor drinking was related to the risk of HAMI TSP.

SHINKAN TOKUDOMO
Department of Public Health,
Nagoya City University Medical School, Nagoya
HIROSHI SHIBASAKI
Department of Brain Pathophysiology,
Faculty of Medicine, Kyoto University, Kyoto
YASUTO ITOYAMA
Department of Internal Medicine,
Faculty of Medicine, Kyushu University, Fukuoka
HIROSHI SHIBASAKI
Department of Internal Medicine,
School of Medicine, Kurume University, Kurume
TATSUOFU NAKAMURA
First Department of Internal Medicine,
Nagasaki University School of Medicine,
Nagasaki
YASUKI MIYOSHI
Department of Neurology,
Kyushu Renai Hospital, Kitakyushu
YOSHIYUKI MURAI
Department of Neurology
MASATO IKEDA
Department of Occupational Health Economics,
University of Occupational and Environmental Health,
Kitakyushu, Japan

Suppression of motor neuron firing by transcranial magnetic stimulation in a patient with multiple sclerosis

Transcranial magnetic stimulation produces a short latency excitatory response in tonically active single motor units in small hand muscles. This is shown by a peristimulus time histogram, which cross correlates motor unit firing times with the time of the stimulus. The excitatory response in the first dorsal interosseous muscle (FDI) occurs at 20–30 ms and is termed the primary peak (PP); in some motor units a secondary peak at 50–90 ms is also seen. PP is characteristically followed by a compensatory period of zero firing probability, which reflects the advancement of discharges that would have occurred shortly after PP had no stimulus been applied. Complete suppression of firing in response to transcranial magnetic stimulation in the absence of an excitatory peak has not been reported. The response of 21 single motor units to transcranial magnetic stimulation has been recorded in nine patients with multiple sclerosis (MS), to study the neural mechanism of symptoms caused by upper motor neuron lesions. Findings have suggested that spatial and temporal summation at the spinal motorneuron is impaired by a reduction in the velocity and synchronisation of central transmission. For healthy subjects, however, complete suppression of tonic activity in single motor units by transcranial magnetic stimulation was not observed.

The patient, a 48 year old female, initially presented in 1986 with paraparesis, weakness affecting the left arm, and diplopia. The diagnosis of MS was supported by the finding of oligoclonal bands in the CSF, enhancing white matter lesions on CT and by delayed VEs. She was subsequently free of symptoms for four years. She then re-presented with pyramidal weakness of the right arm, [reducing the power of the right FDI to 4/5, MRC scale], increased tone in the left leg, extensor plantar responses and impaired joint position sense in the toes. During this clinical episode the response to transcranial magnetic stimulation of three voluntarily activated low threshold motor units from the weakened right FDI was recorded.

The patient was right handed and gave her informed consent to the experiments which were performed with the approval of the local ethics committee. The inducing current flowed in an anticlockwise direction through a circular (Novametrix 200) positioned tangentially at the vertex. Single MU potentials were recorded using fine concentric needle electrodes (Dantec type 13L58). The patient was asked to maintain repetitive motor unit firing and was aided by auditory and visual feedback of the motor unit discharge. Signals were amplified with a band pass of 32 Hz to 16 kHz (Medelec type MS6) and epochs of -250 to +250 ms relative to the stimulus were digitised at 10 kHz (Cambridge Electrleic Design 1401) for subsequent analysis. Up to 120 stimuli were given for each motor unit, delivered at random with respect to the time of the stimulus.
HAM/TSP attributable to blood transfusion.

S Tokudome, H Shibasaki, Y Itoyama, H Shoji, T Nakamura, T Miyoshi, Y Murai and M Ikeda

*J Neurol Neurosurg Psychiatry* 1992 55: 738
doi: 10.1136/jnnp.55.8.738

Updated information and services can be found at:
http://jnnp.bmj.com/content/55/8/738.1.citation

**Email alerting service**

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/